



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵: B01D 71/38, 69/12, C02F 1/44	A1	(11) International Publication Number: WO 94/17905 (43) International Publication Date: 18 August 1994 (18.08.94)
(21) International Application Number: PCT/GB94/00198 (22) International Filing Date: 2 February 1994 (02.02.94) (30) Priority Data: 9302035.2 2 February 1993 (02.02.93) GB 9306761.9 31 March 1993 (31.03.93) GB (71) Applicant (for all designated States except US): NORTH WEST WATER GROUP PLC. [GB/GB]; Dawson House, Great Sankey, Warrington WA5 3LW (GB). (72) Inventors; and (75) Inventors/Applicants (for US only): COLQUHOUN, Howard, Matthew [GB/GB]; The Field, Goughs Lane, Knutsford, Cheshire WA16 8QN (GB). WILLIAMS, Peter, John [GB/GB]; 7 Ennerdale Drive, Frodsham, Cheshire WA6 7LS (GB). LEWIS, Andrew, Lennard [GB/GB]; 1 Lllys Derw, School Lane, Ponciau, Wrexham, Clwyd LL14 1RP (GB). (74) Agent: ATKINSON, Peter, Birch; Marks & Clerk, Suite 301, Sunlight House, Quay Street, Manchester M3 3JY (GB).		(81) Designated States: AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, ES, FI, GB, HU, JP, KP, KR, KZ, LK, LU, LV, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SK, UA, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: POLYMER POROUS STRUCTURE AND PROCESS		
(57) Abstract <p>A reverse osmosis membrane comprises a support membrane which itself has properties of salt rejection laminated to a layer comprising a cross-linked optionally protected polyol. The support membrane may be a reverse osmosis membrane and the polyol may be polyvinyl alcohol. The membrane of the invention may be prepared by the following steps: a) coating the surface of a support membrane which itself has properties of salt rejection with a solution comprising a polyol; b) cross-linking the polyol, and c) as necessary polymerising and/or protecting the polyol.</p>		

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POLYMER POROUS STRUCTURE AND PROCESS

This invention relates to a novel class of polymer membranes, particularly to a structure having high flux and salt rejection, which can be used to advantage in reverse osmosis, and to a process for the construction of such a membrane.

Reverse osmosis membranes, whether integral asymmetric or composite in structure, are known, e.g. those comprising a layer of a sulphonated polysulphone active material, and the latter often have an acceptable performance in terms of water flux and rejection of dissolved salts, the active material providing the good salt rejection and having good chemical stability, e.g. chlorine-, pH- and heat-resistance. However, in general, both the flux and salt rejection of such membranes, represent a compromise level of performance, which is no more than acceptable. Thus, if the water flux of such membranes is at a commercially viable level of performance, the salt rejection is at a level of performance which is merely 'average' and vice versa. This means that such membranes can in practice only be used in the reverse osmosis (i.e. in e.g. desalination) of feedwaters with a relatively low salt content, or with the additional cost of recycling any feedwater with a higher salt content.

There is clearly a need for a reverse osmosis membrane with a structure having high flux and an active material having good salt rejection and chemical stability, the performance of which in both aspects is appreciably better than that of known membranes which have good chemical stability, e.g. those comprising a layer of a sulphonated polysulphone.

By 'appreciably better' herein we mean that the salt transmission is less than 40% of the value for the corresponding known membrane, whilst the water flux is in excess of 40% of the value for the corresponding known membrane. Preferably, the salt transmission is less than 30%, in particular less than 20%, of the value for the

corresponding known membrane, whilst the water flux is preferably in excess of 50%, in particular more than 60%, of the value for the known membrane.

We have now obtained high performance reverse osmosis membranes of this type, in which we have for the first time been able to select a flux level also providing good salt rejection.

Accordingly, the present invention provides a reverse osmosis membrane comprising a support membrane which itself has properties of salt rejection laminated to a layer comprising a cross-linked optionally protected polyol.

Such membranes may be used in the reverse osmosis (i.e. in e.g. desalination) of feedwaters with a relatively high salt content, with minimisation of the cost of recycling any such feedwater with a higher salt content.

The support membrane is one which (in the absence of the layer of cross-linked optionally protected polyol) has salt rejection properties. This support membrane is preferably a reverse osmosis membrane but may be a nanofiltration membrane.

The membranes of the invention may be classed as composite membranes by virtue of their construction which incorporates the support membrane (which itself has salt rejection properties) and the cross-linked polyol layer laminated thereto. The support membrane may, for example, be an integral asymmetric membrane. It is however more preferred that the support membrane is itself a composite membrane (referred to herein as a composite support membrane to distinguish it from the membranes of the invention). The composite support membrane may for example comprise a layer providing a degree of salt rejection laminated to an ultrafiltration support membrane. Membranes in accordance with the invention comprising a composite support membrane and a layer of cross-linked polyol are referred to herein as multi-layer composite membranes.

Preferably the polyol has a weight average molecular

weight of 500 to 500,000.

The polyol in any membrane of the present invention will often be cross-linked by means of an active cross-linking agent.

The cross-linking agent may be any species which is capable of reacting with the polyol hydroxyl functions intermolecularly, e.g. an organic compound having a reactive carbonyl group or a precursor thereof which is capable of so reacting. Such compounds include aliphatic aldehydes and ketones, e.g. formaldehyde or butyraldehyde, which react with the polyol hydroxyl functions, in the presence of a strong acid, by acetal formation.

The polyol in the membrane of the present invention is thus believed (without prejudice to the present invention as claimed) in such cases to be cross-linked by hydrocarbylenedioxy groups.

The polyol in any membrane of the present invention will often be protected conventionally. The protecting agent may be any species which is capable of reacting with the polyol hydroxyl functions intramolecularly, e.g. an organic compound having a reactive group, e.g. a carbonyl group or a precursor thereof which is capable of so reacting. Such compounds include oxoalkanes, such as aliphatic aldehydes and ketones, e.g. formaldehyde or butyraldehyde which can react with the polyol, in the presence of a strong acid, by acetal formation.

The polyol in the membrane of the present invention is thus believed (without prejudice to the present invention as claimed) in such cases to be protected in part by hydrocarbylenedioxy groups replacing pairs of hydroxyl functions.

From the foregoing, it will be seen that the polyol component of the present membrane may be cross-linked and optionally protected by a single agent for cross-linking and protecting the polyol, which may be one having a reactive carbonyl group or a precursor thereof which is capable of reacting with and linking pairs of hydroxyl

functions.

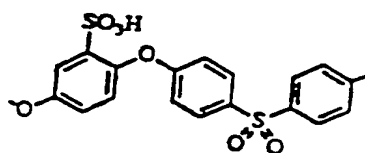
All materials for the support membrane should of course

- a) have adequate resistance to their chemical environment in the process in which the reverse osmosis membrane of the invention is to be used, and
- b) not be prone to fouling in that environment.

Any material may be used in the support membrane of the present invention provided that it has an adequate stability for its intended use; preferably it has a high chemical stability, so that its performance is not appreciably affected by the nature of any impurities, such as chlorine, acid or alkali, in, or the temperature of, the feedwater.

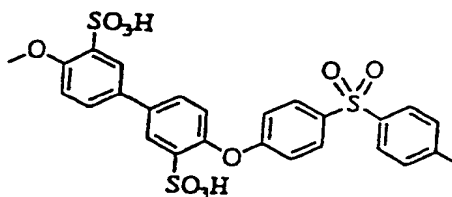
Materials for such support membranes include those known as SPEES/PES (i.e. sulphonated poly(ether ether sulphone)/poly(ether sulphone)) and SPEDES/PES (i.e. sulphonated poly(ether diphenyl ether sulphone)/poly(ether sulphone)). SPEES/PES and SPEDES/PES are each based respectively on a copolymer of a unit of formula I or II:

SEES



I

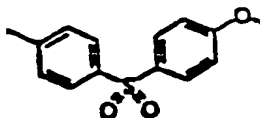
SEDES



II

with a unit represented by the following formula III:

ES



III

if the polymer is SPEDES/PES then the preferred molar ratio range of ES:SEDES is 1:1 to 20:1, e.g. 2:1 to 12:1 (e.g. 2:1 to 10:1), in particular 3:1 to 8:1.

If the polymer is SPEES/PES then the preferred molar ratio range for ES:SEES is 1:2 to 10:1 (in particular 1:1 to 5:1).

Each such sulphonated unit is hereinafter referred to for convenience by the trivial name appended to it above.

These materials have a good chemical stability, e.g. chlorine-, pH- and heat-resistance, and are also advantageous in other aspects.

These are especially advantageous in that the initial unsulphonated polymeric precursor of the sulphonated polymer is easily obtained by polycondensation of hydroquinone or 4,4'-biphenol, 4,4'-dihydroxydiphenylsulphone and 4'4'-dichlorodiphenylsulphone.

These polymeric precursors are also advantageous in that they are rapidly, selectively and quantitatively sulphonated only on the dioxydiphenylene or dioxyphenylene residue at the positions shown with no degree of molecular weight degradation during sulphonation.

The degree of sulphonation, and hence the level of hydrophilicity, of the final polymer is readily predetermined by the molar ratio of the SEDES or SEES to ES.

Any SPEDES/PES and SPEES/PES material which has a composition within the ranges given hereinbefore, and can give the required good salt rejection may be used in the support layer of the reverse osmosis membranes of the

present invention.

Preferred reverse osmosis membranes of the present invention have, however, in addition to high chemical stability, a performance which has not only appreciably better as defined hereinbefore, but exhibit the levels of performance values identified as preferred.

These properties are largely determined by the composition of the SPEDES/PES or SPEES/PES polymer.

Accordingly, in a preferred embodiment, the present invention provides a reverse osmosis membrane in which the active layer (i.e. the support layer) comprises a SPEDES/PES polymer, in which the molar ratio of the SEDES and ES units are in the ranges of ES:SEDES of 2:1 to 12:1, in particular 3:1 to 8:1.

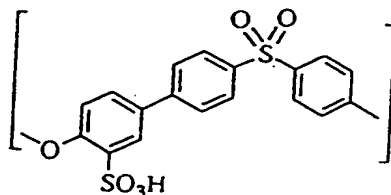
Alternatively the support layer may comprise a SPEES/PES polymer in which the molar ratio of the SEES and ES units are in the ranges of ES:SEES 1:2 to 10:1, in particular 1:1 to 5:1.

These anionic materials of the active layer of the present membrane will of course contain a counterion.

This may be any cation, but is usually a hydrogen cation, or an alkali metal cation, such as sodium or potassium.

Further examples of polymers which may be used as the active layer are those incorporating sub-units of the formula (IV)

SEDS



(IV)

either alone or in combination with units of the formula (III) (see above)

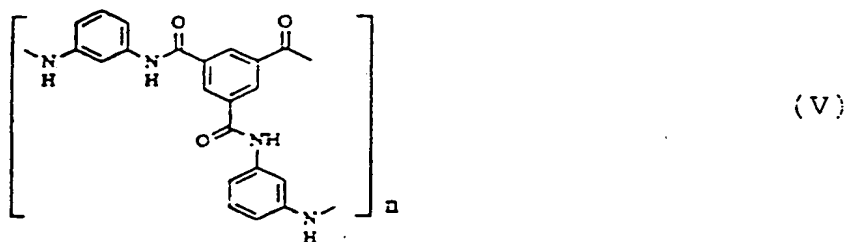
Such polymers may be prepared by polycondensation or

copolycondensation (as the case may be) of 4-(4-chlorophenylsulphonyl)-4'-hydroxydiphenyl for which a synthesis is described in Polymer 1977, Vol 18, April, pages 354 et seq. The resultant polymer is then sulphonated. Copolymers containing units (IV) and (III) are referred to herein as SPEDS/PES.

In the membrane of the present invention, other suitable materials for the support (i.e. active) layer include polyamides, such as unsaturated aromatic polyamides and araliphatic polyamides, polyureas, polyimides, such as unsaturated aromatic and araliphatic polyimides, polyaroazoles and polyarodiazoles, such as polybenzimidazole, polyhydrazides, polyesters, cellulosics, such as cellulose diacetate and triacetate, and anion-substituted polyacrylics and polyvinyls.

Accordingly, in another embodiment the present invention provides a reverse osmosis membrane in which the support (active) layer comprises a cross-linked polyamide, polyamide, polyaroazole or polyarodiazole which is laminated to the layer of cross-linked optionally protected polyol.

One class of such active materials in the membrane include a cross-linked aromatic polyamide approximately represented by the following formula V:



When the present membrane is a multi-layer composite membrane, it will include a composite support membrane. This composite support membrane will comprise a layer which itself has a degree of salt rejection, laminated to a substrate membrane (e.g. an ultrafiltration membrane).

Thus a suitable composite support membrane will include a substrate membrane comprising

- a) a polymer, such as a polyarylsulphone, a polyarylethersulphone, polyvinylidene fluoride or a polyaryletherketone, or
 - b) a ceramic, such as zirconia,
- the latter being less preferred on the grounds of cost.

Where, as is preferred a polymer, such as a polyarylsulphone, a polyarylethersulphone, a poly(ether-imide), a polyvinylidene fluoride or a polyaryletherketone, is used as the substrate membrane, the support membrane will often also comprise a reinforcement of a material which is also not appreciably affected by the nature of any impurities, such as chlorine. acid or alkali, in, or the temperature of, the feedwater.

It may be e.g. a conventional cloth backing sheet of a material such as polypropylene or polyester with high chemical resistance, on which the polymer of the substrate membrane, such as a polyarylsulphone, a polyarylethersulphone, polyvinylidene fluoride or a polyaryletherketone, is present as a coating.

Polysulphone-based substrate membranes are particularly preferred for their high chemical resistance, e.g. those of the polysulphone UDEL or the polyarylethersulphone PES (see below).

The substrate membrane should preferably have an effective pore size in the range of 50Å to 50,000Å, in particular 500Å to 5000Å.

Such substrate membranes should have a pure water permeability of 0.033 to 3.3 m/day at 1 bar, e.g. 0.1 to 1.0 m/day at 1 bar.

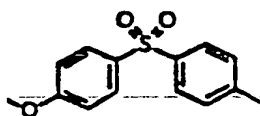
Where the multilayer composite membrane of the invention also comprises a reinforcement in the form of a backing sheet which bears a layer of a polymer, such as a polyarylsulphone, a polyarylethersulphone, a poly(ether-imide), a polyvinylidene fluoride or a polyaryletherketone, the latter will often have a thickness in the range of 20

μ to 250 μ , in particular 30 μ to 100 μ .

Multi-layer composite membranes including a backing sheet should have a total thickness in the ranges of 50 μ to 500 μ , in particular 100 μ to 250 μ .

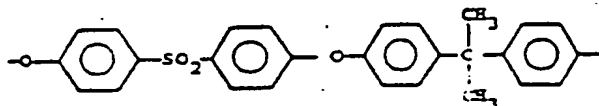
As examples of polymers of which the substrate membrane may be made are inter alia polymers and copolymers of units of the general formulae III and VI-XII.

ES



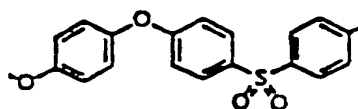
III

UDEL



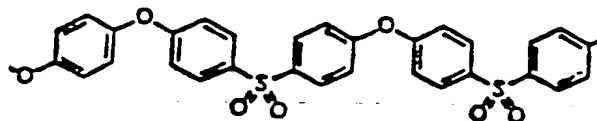
VII

EES



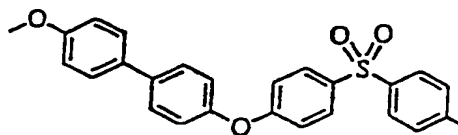
VIII

EESES



IX

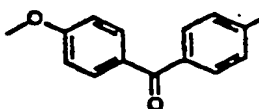
EDES



X

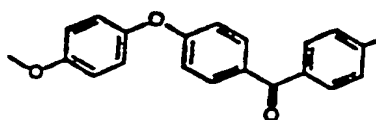
The polymer is preferably a homopolymer e.g. UDEL or PES.

However, we do not exclude the possibility that it may be a copolymer e.g. PEES/PES, or a polyetherketone such as one wherein the comonomer units are represented by the following formulae:



EK

XI



EEK

XII

Each such polymer type is hereinafter referred to for convenience by the trivial name appended to it above.

The present invention also provides a process for the preparation of a reverse osmosis membrane, comprising the steps of

- a) coating a surface of a membrane which itself has salt rejection properties or a component layer thereof with a solution comprising a polyol,
- b) cross-linking the polyol, and
- c) as necessary polymerising and/or protecting the polyol.

In a preferred embodiment,

- a1) the polyol in the coating solution is polymeric, and
- b1) cross-linking the polyol and protection thereof as necessary, is effected by a single component of the coating solution.

In step a1, the polymeric polyol may be a polymer having repeating units formally derived from a polymerised unsaturated lower alcohol. The preferred polyol is polyvinyl alcohol.

The single component of the coating solution for cross-linking and optionally protecting the polyol may be an organic compound having a reactive carbonyl group or a precursor thereof which is capable of reacting with the polyol alcohol functions.

Compounds having a reactive carbonyl group or a precursor thereof which is capable of reacting in this way include aliphatic aldehydes and ketones, e.g. formaldehyde or butyraldehyde, in the presence of a strong acid.

Where the compound which has a group or a precursor thereof which is capable of reacting with the polyol alcohol functions is an aldehyde or ketone in the presence of a strong acid, the strong acid may be sulphuric acid, phosphoric acid or a sulphonic acid.

The support membrane is preferably a reverse osmosis membrane but may for example be a nanofiltration membrane. If the support membrane is itself a composite support membrane then the process of the invention comprises coating the surface of the active layer thereof.

Typically, the coating solution is applied as a thin film and the solvent is subsequently removed.

In the preferred embodiment, this is typically effected at a temperature between 20 and 80°C, which, it is believed, simultaneously cross-links some of the polyol alcohol functions intermolecularly and links and protects some of the polyol alcohol functions intramolecularly, in both cases by acetal formation, giving rise to a reverse osmosis membrane of the present invention.

The solvent in the coating solution, which is removed after coating, should (especially in the preferred embodiment) be chosen so that the surface of the reverse osmosis (optionally composite) membrane which is coated is not unduly attacked or swollen (and preferably is

unaffected) by the solvent at the temperature at which

- a) the surface is coated and/or
- b) the solvent is removed.

The solvent may thus often conveniently consist mainly of water and/or a lower alcohol, provided that the same is compatible with any agent e.g. organic compound for cross-linking and optionally protecting the polyol functional groups in the product membrane.

The present invention will now be illustrated by the following Examples:

Example 1

A composite reverse osmosis membrane comprising a selective layer of a SPEES/ES (1:3 molar ratio) copolymer (i.e. of a monomer of formula I with a comonomer of the formula III in the molar ratio of 1:3), laminated to a polysulphone ultrafiltration support-membrane was coated with a 1 wt% aqueous solution of polyvinyl alcohol (65,000 MW (weight average), 100% hydrolysed) containing 1 wt% sulphuric acid and 4 wt% formaldehyde using a continuous bead-coater.

The resulting coated membrane was dried at 40°C and tested in reverse osmosis using 0.2% sodium chloride solution at 40 bar pressure.

The results were as follows:

	<u>Flux (GFD)</u>	<u>Salt Rejection %</u>	<u>Salt Transmission%</u>
Before	49.3	96.4	3.6
After	36.2	99.1	0.9
% Original	73.4%		25.0%

Example 2

A composite reverse osmosis membrane comprising a selective layer of SPEDES/ES (1:6 molar ratio) copolymer

(i.e. of a monomer of formula II with a comonomer of the formula III in the molar ratio of 1:6), laminated to a polysulphone ultrafiltration support-membrane was coated with a 1wt% aqueous solution of polyvinyl alcohol (65,000 MW (weight average), 100% hydrolysed) containing 1 wt% sulphuric acid and 4 wt% formaldehyde using a continuous bead-coater.

The resulting coated membrane was dried at 40°C and tested in reverse osmosis using 0.2% sodium chloride solution at 40 bar pressure.

The results were as follows:

	<u>Flux (GFD)</u>	<u>Salt Rejection %</u>	<u>Salt Transmission%</u>
Before	104.0	89.4	10.6
After	64.0	97.0	3.0
% Original	62.0%		28.0%

Example 3

A commercially available composite reverse osmosis membrane (Filmtec FT-30), comprising a selective layer of polymer of the formula V laminated to a polysulphone ultrafiltration support-membrane, was coated with a 1 wt% aqueous solution of polyvinyl alcohol (65,000 MW (weight average), 100% hydrolysed) containing 1 wt% sulphuric acid and 4 wt% formaldehyde using a continuous bead-coater.

The resulting coated membrane was dried at 40°C and tested in reverse osmosis using 0.2% sodium chloride solution at 40 bar pressure.

The results were as follows:

	<u>Flux (GFD)</u>	<u>Salt Rejection %</u>	<u>Salt Transmission%</u>
Before	69.4	97.6	2.4
After	39.1	99.6	0.4
% Original	56.3%		16.7%

Example 4

A composite reverse osmosis membrane comprising a selective layer of SPEDS/PES (1:3 mole ratio), laminated to a polysulphone ultrafiltration support membrane was coated with a 1 wt% aqueous solution of polyvinyl alcohol (65,000 MW (weight average), 100% hydrolysed) containing 1 wt% sulphuric acid and 4 wt% formaldehyde using a continuous bead-coater. The resulting coated membrane was dried at 50°C and tested in reverse osmosis using 0.2% sodium chloride solution at 40 bar pressure. The results before and after coating were as follows:-

	<u>Flux (GFD)</u>	<u>Salt Rejection %</u>	<u>Salt Transmission%</u>
Before	62.2	95.9	4.1
After	26.7	99.5	0.5
% Original	42.9%		12.2%

CLAIMS

1. A reverse osmosis membrane comprising a support membrane which itself has properties of salt rejection laminated to a layer comprising a cross-linked optionally protected polyol.

2. A membrane as claimed in claim 1 wherein the polyol has been cross-linked by an organic compound having a reactive carbonyl group or a precursor thereof which is capable of so reacting.

3. A membrane as claimed in claim 2 wherein the polyol has been cross-linked by a aliphatic aldehyde or ketone.

4. A membrane as claimed in claim 3 wherein the polyol has been cross-linked by formaldehyde or butyraldehyde.

5. A membrane as claimed in claim 2 wherein the polyol has been cross-linked by hydrocarbylenedioxy groups.

6. A membrane as claimed in any one of claims 1 to 5 wherein the polyol is protected.

7. A membrane as claimed in claim 6 wherein the polyol has been protected by intramolecular reaction of hydroxyl groups with the compound used for effecting cross-linking of the polyol.

8. A membrane as claimed in any one of claims 1 to 7 wherein the polyol has a weight average molecular weight of 500 to 500,000.

9. A membrane as claimed in any one of claims 1 to 8 wherein the polyol is a polymer having repeating units formally derived from a polymerised lower unsaturated alcohol.

10. A membrane as claimed in any one of claims 1 to 8 wherein the polyol is polyvinyl alcohol.

11. A membrane as claimed in any one of claims 1 to 10 wherein the support membrane comprises SPEES/PES or SPEDES/PES.

12. A membrane as claimed in claim 11 wherein the support membrane comprises SPEDES/PES and the molar ratio of ES:SEDES is in the range 1:1 to 20:1.

13. A membrane as claimed in claim 12 wherein said molar ratio range is 2:1 to 12:1.

14. A membrane as claimed in claim 13 wherein said molar ratio range is 3:1 to 8:1.

15. A membrane as claimed in claim 11 wherein the support membrane comprises SPEES/PES and the molar ratio of ES:SEES is in the range 1:2 to 10:1.

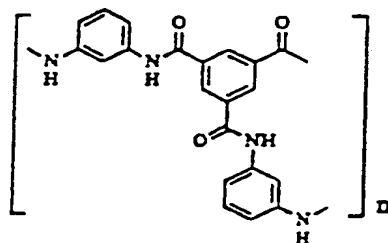
16. A membrane as claimed in claim 15 wherein said molar ratio range is 1:1 to 5:1.

17. A membrane as claimed in any one of claims 1 to 10 wherein said support membrane comprises SPEDS or SPEDS/PES.

18. A membrane as claimed in any one of claims 1 to 10 wherein said support membrane comprises a polyamide, polyurea, polyimide, polyaroazole, polyarodiazole, polyhydrazide, polyester, cellulosic, or an anion-substituted polyacrylic or polyvinyllic.

19. A membrane as claimed in claim 18 wherein the support membrane comprises a cross-linked polyamide, polyimide, polyaroazole, or polyarodiazole.

20. A membrane as claimed in claim 19 wherein the support membrane comprises a cross-linked aromatic polyamide approximately represented by the following formula V.



(V)

21. A multi-layer composite membrane comprising a reverse osmosis membrane as claimed in any one of claims 1 to 20 associated with a porous substrate membrane.

22. A multi-layer composite membrane as claimed in claim 21 wherein the substrate membrane comprises a polymer or a ceramic.

23. A multi-layer composite membrane as claimed in claim 22 wherein the substrate membrane is a polyarylsulphone, a polyarylethersulphone, a poly(ether-imide), a polyvinylidene fluoride or a polyaryletherketone.

24. A multi-layer composite membrane as claimed in claim 23 wherein the substrate membrane is of zirconia.

25. A multi-layer composite membrane as claimed in any one of claims 21 to 24 wherein the substrate membrane has a pore size in the range 50Å to 50,000Å.

26. A multi-layer composite membrane as claimed in claim 25 wherein the substrate membrane has a pore size in the range 500Å to 5000Å.

27. A multi-layer composite membrane as claimed in any one of claims 21 to 26 having a total thickness in the range 50 μ to 500 μ .

28. A multi-layer composite membrane as claimed in claim 27 having a total thickness of 100 μ to 250 μ .

29. A process for preparing a reverse osmosis membrane comprising the following steps

- a) coating the surface of a support membrane which itself has properties of salt rejection with a solution comprising a polyol,
- b) cross-linking the polyol, and
- c) as necessary polymerising and/or protecting the polyol.

30. A process as claimed in claim 29 wherein cross-linking and protection of the polyol is effected by a single component of the coating solution.

31. A process as claimed in claim 30 wherein said component is an organic compound having a reactive carbonyl group or precursor thereof.

32. A process as claimed in claim 31 wherein said component is an aliphatic aldehyde or ketone and said cross-linking/protection reaction is effected in the presence of a strong acid.

33. A process as claimed in claim 32 wherein said component is formaldehyde or butyraldehyde.

34. A process as claimed in any one of claims 29 wherein the polyol is a polymer having repeated units formally derived from a polymerised lower unsaturated alcohol.

35. A process as claimed in claim 34 wherein said polyol is polyvinyl alcohol.

36. A process as claimed in any one of claims 30 to 35 wherein the solvent for the coating solution comprises water and/or a lower alcohol.

37. A reverse osmosis process characterised by the use of a membrane as claimed in any one of claims 1 to 28.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/GB 94/00198A. CLASSIFICATION OF SUBJECT MATTER
IPC 5 B01D71/38 B01D69/12 C02F1/44

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 5 B01D C02F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	DE,A,38 01 528 (ZWECKVERBAND SONDERMÜLLPLÄTZE MITTELFRANKEN) 27 July 1989 see the whole document ----	1
A	EP,A,0 047 953 (ALIGENA AG) 24 March 1982 see the whole document ----	1
A	EP,A,0 442 557 (METALLGESELLSCHAFT AG) 21 August 1991 ----	1
A	EP,A,0 228 248 (NITTO ELECTRIC INDUSTRIAL CO., LTD.) 8 July 1987 ----	1
A	FR,A,2 371 477 (UOP INC.) 16 June 1978 -----	1

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

26 May 1994

Date of mailing of the international search report

07.06.94

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INTERNATIONAL SEARCH REPORT

information on patent family members

International application No.

PCT/GB 94/00198

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		US-A- 5183571	02-02-93
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FR-A-2371477	16-06-78	NONE	